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LISTING OF CLAIMS:

17. (currently amended) A process for minimizing thermal aggregation of DNase in a liquid solution comprising:

introducing a DNase aggregation-inhibiting amount of sugar to a solution comprising a <u>human DNase</u>, wherein and elevating the temperature of said DNase solution is subsequently <u>elevated to above 37°C and aggregation of DNase at said elevated</u> temperature is reduced.

- 18. (previously presented) A process according to claim 17, wherein the temperature of said solution is elevated above about 60°C.
- 19. (currently amended) A process according to claim 17, further-comprising reducing wherein the pH of said solution is below pH 7.0.
- 20. (previously presented) A process according to claim 19, wherein said solution is at about pH 6.5.
- 21. (previously presented) A process according to claim 19, wherein said solution is at about pH 6.
- 22. (previously presented) A process according to claim 19, wherein said solution is at about pH 5.
- 23. (previously presented) A process according to claim 17, wherein said sugar is present in an amount from 50 mg/ml to 200 mg/ml.
- 24. (currently amended) A process according to claim 17, wherein said sugar is selected from the group consisting α -lactose monohydrate, mannitol, trehalose or and sucrose-
- 25. (previously presented) The process according to claim 17, further comprising the steps of spray-drying said liquid solution and collecting the spray-dried product as a respirable

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DNase-containing powder that is therapeutically effective when administered into the lung of an individual.

- 26. (currently amended) A DNase solution comprising <u>a human DNase</u> and a DNase aggregation-inhibiting amount of sugar wherein said DNase solution is minimally aggregated when said solution is at a temperature greater than 37°C.
- 27. (previously presented) A DNase solution according to claim 26, wherein the temperature is greater than about 60°C.
- 28. (currently amended) A DNase solution according to claim 26, wherein the pH of said solution is further kept at a pH below 7.0.
- 29. (previously presented) A DNase solution according to claim 28, wherein said solution is at about pH 6.5.
- 30. (previously presented) A DNase solution according to claim 28, wherein said solution is at about pH 6.
- 31. (previously presented) A DNase solution according to claim 26, wherein said sugar is present in an amount from 50 mg/ml to 200 mg/ml.
- 32. (currently amended) A DNase solution according to claim 26, wherein said sugar is selected from the group consisting α -lactose monohydrate, mannitol, trehalose of and sucrose.
- 33. (currently amended) A composition comprising the A DNase solution according to claim 26, wherein said solution that is further spray-dried to a respirable DNase-containing powder that is therapeutically effective when administered into the lung of an individual.